Detection of CXCL12/SDF-1 in Formalin Fixed Paraffin-Embedded Human Tissue

Reagents:

1X Automation Buffer
3% Hydrogen Peroxide
Antibody Diluent
Citrate Buffer
DAB Chromagen
Hematoxylin

Antibody Information

Blocking Serum: Normal Horse Serum
Jackson Immunoresearch Laboratories, Inc.
West Grove, PA 19390
www.jacksonimmuno.com
1-800-367-5296
Catalog # 008-000-001

Avidin/Biotin Blocking Kit Vector Laboratories, Inc. Burlingame, CA 94010 www.vectorlabs.com 1-800-227-6666 Catalog # SP-2001

Primary antibody: Monoclonal anti-human/mouse CXCL12/SDF-1 Antibody R&D Systems
Minneapolis, MN 55413
www.rndsystems.com
1-800-343-7475
Catalog # MAB350

Negative control: Normal Mouse Serum
Jackson Immunoresearch Laboratories, Inc.
West Grove, PA 19390
www.jacksonimmuno.com
1-800-367-5296
Catalog # 015-000-001

Secondary antibody: Biotinylated horse anti-mouse IgG Vector Laboratories, Inc.
Burlingame, CA 94010
www.vectorlabs.com

1-800-227-6666 Catalog # BA2001

<u>Label antibody: Peroxidase-conjugated Streptavidin SS Label</u>

Biogenex

San Ramon, CA 94583

www.biogenex.com

1-800-421-4149

Catalog # HK330-9K

Staining Procedure

Positive Control Tissue: Tonsil

Stain Localization: Cytoplasmic - endothelial cells and crypts

Deparaffinize and hydrate slides through the following solutions.

Xylene	2 times	5 minutes
100% Ethanol	2 times	3 minutes
95% Ethanol	2 times	3 minutes
1X Automation Buffer	2 times	5 minutes

- 1. Quench endogenous peroxidase by placing slides in 3% hydrogen peroxide for 15 minutes.
- 2. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.

2. Killse slides ili 2 e	nanges of 1A Automatio	ii Duiici ioi 5 iiiiiuu	es caen.
3. Unmasking Techn	iques using the decloake	r.	
Add 500 ml of dis	tilled water to the pan of	the decloaker.	
Place full rack of s	slides in 200 ml of 1X cit	rate buffer and place	in the decloaker
Decloak for 5 min	. Pressure		
Depressurize for 1	0 minutes.		
Remove pan top a	nd cool for 10 minutes. T	Temperature after co	oling
Rinse in distilled v	vater two times for 3 mir	nutes each.	
4. Rinse slides in 2 c	hanges of 1X Automation	n Buffer for 5 minute	es each.
5. Block with 10% N	formal Horse Serum for 2	20 minutes at room to	emperature.
Lot#	Reconstituted Date	<u> </u>	_
DO NOT DINCE CI	IDES. CONTINUE TO	AVIDIN DIOTIN D	LOCK
DO NOT KINSE SL	IDES. CONTINUE TO	AVIDIN-DIOTIN D	LUCK.
6. Apply the Avidin	Biotin Blocking kit		
11 .	Exp Date	New Kit:	yes / no

Apply avidin block - 15 minutes at room temperature. Quick rinse in 1X Automation Buffer. Apply biotin block - 15 minutes at room temperature. Wipe excess block.

DO NOT RINSE SLIDES WITH BUFFER BEFORE ADDING PRIMARY ANTIBODY.

7. Apply primary antibody (CXCL12) at a 1:150 dilution and incubate for one hour at room temperature.	
Lot#Exp Date	
For negative control slides, normalize the protein concentration of the normal mouse serum to the protein concentration of the primary antibody (CXCL12), and use this t make a 1:150 dilution. Apply to the slides and incubate for one hour at room temperature. Lot# Reconstituted Date	
8. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.	
9. Apply biotinylated horse anti-mouse secondary at 1:1000 and incubate for 30 minute at room temperature. Lot#	38
10. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.	
11. Apply Biogenex SS Label and incubate for 30 minutes at room temperature. Lot# Expiration Date	
12. Rinse slides in 2 changes of 1X Automation Buffer for 5 minutes each.	
13. Apply liquid Dako DAB Chromagen for 6 minutes in the dark. (Add 1 drop of DAB per ml of substrate) Lot#	
14. Rinse in tap water 3 minutes.	
15. Counterstain with Modified Harris Hematoxylin for 20 seconds.	
16. Rinse in tap water until water is clear.	
17. Gently agitate slides in 1X Automation Buffer until blue.	
18. Dehydrate through the following solutions.	

95% Ethanol	1 change	3 minutes
100% Ethanol	3 changes	3 minutes
Xylene	2 changes	5 minutes

19. Coverslip

Updated 08/21/06